AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

- 1. (Currently Amended) A method of creating a nucleic acid multiplex, said method comprising the steps of:
- 1) creating a mixture comprising water, a Watson-Crick duplex, a sufficient number of single-stranded mixed base sequence molecules to form a-the multiplex that includes including the Watson-Crick duplex, and an accelerator agent that increases a rate or amount of multiplex formation, said multiplex being a triplex or quadruplex, wherein said single stranded molecule or molecules are selected so that, if in a multiplex, they would each be related to all other strands of the multiplex by adherence to base pairing rules, said rules being either Watson Crick base pairing rules or homologous binding base pairing rules; and
- 2) incubating said mixture to allow the multiplex to form, each strand of said multiplex related to all other strands of the multiplex by adherence to <u>Watson-Crick base-pairing rules or</u> homologous binding base-pairing rules;

provided that, within the multiplex, the Watson-Crick duplex added in step (1) is heteropolymeric with a G-C content between 10% and 90% and a combined frequency therein of purine-pyrimidine dimers and pyrimidine-purine dimers exceeds 25%.

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2. (Original) A method of Claim 1 wherein the multiplex created is a triplex, in step (1) the sufficient number of single-stranded molecules is 1, and in step (2) the triplex is formed.

3. (Original) A method of Claim 1 wherein the duplex substantially retains its double-helical structure and the single-stranded molecule resides in a groove of that double-helical structure.

4. (Currently Amended) A method of Claim 3-1 wherein the single-stranded molecule is related to one strand of the duplex by Watson-Crick base-pairing rules and to the second strand of the duplex by homologous binding base-pairing rules.

5. (Original) A method of Claim 4 wherein the duplex substantially retains its doublehelical structure and the single-stranded molecule resides in a groove of that double-helical structure.

6. (Canceled) A method of Claim 1 where, within the multiplex, the Watson-Crick duplex added in step (1) is heteropolymeric with a G-C content between 10% and 90%, and furthermore the combined frequencies therein of purine-pyrimidine dimers and pyrimidine-purine dimers exceeds 25%.

- 7. (Currently Amended) A method of Claim 1 wherein steps (1) and (2) are performed with at least one of the nucleic acid strands and/or the duplexes not in a cell.
- 8. (Currently Amended) A method of Claim 1 wherein step (2) is performed without the assistance of a protein.

9. (Currently Amended) A method of Claim 1 wherein in step (1), the water is added so

that it accounts, on a volume basis, for at least 50 percent of the <u>a</u> final volume of the mixture.

- 10. (Currently Amended) A method of Claim 1 wherein in step (1), the water is added so that it accounts, on a volume basis, for at least 80 percent of the a final volume of the mixture.
- 11. (Original) A method of Claim 1 wherein in step (1), the water is added so that it accounts, on a volume basis, for all of the liquid added to the mixture.
- 12. (Currently Amended) A method of Claim 1 wherein step (2) is performed at a temperature or temperatures above the <u>a</u> freezing temperature of the <u>aqueous solution mixture</u> and at not more than 85°C.
- 13. (Currently Amended) A method of Claim 12 wherein step (2) is performed at athe temperature or temperatures is/are between 5 °C and to 30 °C.
- 14. (Currently Amended) A method of Claim 13 wherein step (2) is performed at athe temperature or temperatures is/are between 15 °C and to 25 °C.
- 15. (Original) A method of Claim 1 wherein in step (1), a cation is added as the accelerator agent.
- 16. (Original) A method of Claim 15 wherein said cation is Na⁺ provided at a concentration of 50mM to 125mM.
 - 17. (Original) A method of Claim 15 wherein said cation is selected from the group

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consisting of Mn⁺² provided at a concentration of 10mM to 45mM, Mg⁺² provided at a concentration of 10mM to 45mM, and Ni⁺² provided at a concentration of 20mM.

- 18. (Original) A method of Claim 1 wherein in step (1) an intercalator is added as an accelerator agent.
 - 19. (Original) A method of Claim 18 wherein the intercalator is a fluorescent intercalator.
- 20. (Original) A method of Claim 19 wherein the fluorescent intercalator is selected from the group consisting of YOYO-1, TOTO-1, YOYO-3, TOTO-3, POPO-1, BOBO-1, POPO-3, BOBO-3, LOLO-1, JOJO-1, cyanine dimers, YO-PRO-1, TO-PRO-1, YO-PRO-3, TO-PRO-3, TO-PRO-5, PO-PRO-1, BO-PRO-1, PO-PRO-3, BO-PRO-3, LO-PRO-1, JO-PRO-1, cyanine monomers, ethidium bromide, ethidium homodimer-1, ethidium homodimer-2, ethidium derivatives, acridine, acridine orange, acridine derivatives, ethidium-acridine heterodimer, ethidium monoazide, propidium iodide, SYTO dyes, SYBR Green 1, SYBR dyes, Pico Green, SYTOX dyes, and 7-aminoactinomycin D.
- 21. (Original) The method of Claim 1 wherein the accelerator agent is a non-intercalating fluorophore.
- 22. (Original) A method of Claim 21 wherein the non-intercalating fluorophore is selected from the group consisting of biotin, rhodamine, Alexa dyes, BODIPY dyes, biotin conjugates, thiol-reactive probes, fluorescein and derivatives including but not limited to the caged probes, Oregon Green, Rhodamine Green, QSY dyes.
 - 23. (Currently Amended) A method of Claim 1 wherein in step (1) the accelerator agent

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is an intercalator that binds to at least one of the minor groove and/or the major groove of the Watson-Crick duplex.

- 24. (Original) The method of Claim 1 wherein in step (1) the accelerator agent at 25 °C is a liquid.
- 25. (Original) The method of Claim 24 wherein in step (1) the accelerator agent is an organic liquid soluble in water.
- 26. (Original) The method of Claim 1 wherein in step (1) an accelerator agent that is a condensation agent as regards the Watson-Crick duplex is added.
- 27. (Previously Amended) The method of Claim 1 wherein in step (1) an accelerator agent that is a decondensation agent as regards the Watson-Crick duplex is added.

Claim 28 (Withdrawn).

29. (Currently Amended) A method of Claim 1 wherein the multiplex created is a quadruplex, in step (1) the Watson-Crick duplex is a first Watson-Crick duplex, and in step (1) the sufficient number of single-stranded molecules is 2, those single-stranded molecules are in a second Watson-Crick duplex, and in step (2) the quadruplex is formed from said first and second duplexes. Preferably step (1) is done with the two single-stranded molecules already in the second Watson-Crick duplex.

Claims 30-55 (Withdrawn).